

Surveillance Trends in the In Vitro Antibacterial Activity of Tigecycline in the Years 2004 to 2008 - Tigecycline Evaluation Surveillance Trial (T.E.S.T.): A Global Perspective



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Revised Abstract

Objectives: Tigecycline, the first member of the glycyglycines, was marketed in mid 2005 and has demonstrated success against multiply-resistant species and phenotypes. Due to its chemical structure, resistance to tigecycline is reportedly difficult to produce even in the laboratory. The T.E.S.T. program is an ongoing global surveillance with the first post-marketing prospective report of tigecycline and comparator in vitro activity for the years 2004 through 2008. **Methods:** More than 120,000 clinical isolates were collected from 1016 investigative sites in 53 countries worldwide. MICs were determined by broth microdilution according to CLSI guidelines using identical panels. **Results:** Results are given by year for all pathogens. Summary data for tigecycline and key species are as follows:

Organism	2004	2005	2006	2007	2008
Acinetobacter spp.	0.25	1	0.25	1	0.25
Acinetobacter baumannii	0.5	1	0.5	1	0.5
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Acinetobacter baumannii	0.5	1	0.5	1	0.5
Acinetobacter baumannii	0.5	1	0.5	1	0.5

Conclusions: Tigecycline demonstrated no significant shift in MIC values over three years from its pre-marketing (2005) baseline values. Tigecycline activity was retained even against strains resistant to other antimicrobials, including ESBL-producing, multi-resistant *Acinetobacter* spp., methicillin-resistant *S. aureus*, vancomycin-resistant enterococci, and penicillin-resistant *S. pneumoniae*.

Introduction

Tigecycline (formerly GAR-936) is a member of a new class of antimicrobial agents, the glycyglycines. This synthetic analogue of the tetracyclines exhibits significant antibacterial activity that is both bacteriostatic and, in certain instances, bactericidal with killing activity that is as much as fourfold better than vancomycin and daptomycin [1, 2]. The development of tigecycline is important in that tigecycline and other glycyglycines are active against bacterial strains carrying either or both of the two major forms of tetracycline resistance: efflux and ribosomal protection. Certain substituents at the 9-position of the tetracycline molecule restore activity against bacteria harboring genes encoding either or both efflux and ribosomal protection. A single chemical modification of tigecycline overcomes the two molecularly distinct forms of resistance while maintaining activity against susceptible gram-positive, gram-negative, aerobic, and anaerobic bacteria [3]. Furthermore, resistance to tigecycline is difficult to produce even in the laboratory.

Previous studies have demonstrated excellent in vitro activity for tigecycline against clinical and laboratory strains of gram-positive and -negative bacteria with minimum inhibitory concentrations for the 90th percentile inhibited at or below 2 µg/ml, including difficult to treat methicillin-resistant *Staphylococcus aureus* (MRSA), vancomycin-resistant enterococci (VRE), and extended-spectrum beta-lactamase (ESBL) producing *Enterobacteriaceae* [4-6]. This study was undertaken to document the in vitro activity of tigecycline against significant numbers of clinical pathogens collected from a large geographically diverse population over five years time. This study is part of the ongoing global Tigecycline Evaluation and Surveillance Trials (T.E.S.T.) program.

Materials & Methods

- All isolates were derived from blood, respiratory tract, urine (no more than 25% of all isolates), skin, wound, body fluids, and other defined sources. Only one isolate per patient was accepted into the study. More than 120,000 clinical isolates were collected and tested between 2004 to 2008 from 1016 investigative sites in 53 countries worldwide. Isolates were identified to the species level and tested at each site by the participating laboratory.
- Organism collection, transport, confirmation of organism identification, and development and management of a centralized database, were coordinated by Laboratories International for Microbiology Studies (LIMS), a division of International Health Management Associates, Inc. located in Schaumburg, IL, USA.
- All organisms were deemed clinically significant by local participant criteria. Isolate inclusion was independent of medical history, antimicrobial use, age, or gender. All sites identified each study isolate utilizing local laboratory criteria.
- Minimum inhibitory concentrations (MICs) were determined by the CLSI recommended broth microdilution testing method [7]. Tigecycline was supplied by Wyeth Pharmaceuticals (Collegeville, PA, USA). All other agents were supplied by the panel manufacturer, MicroScan (Dade Behring Inc., Sacramento, CA, USA). The following antimicrobial agents were included on the panels with their dilution ranges (expressed in µg/ml): amikacin (0.5-64); amoxicillin/clavulanic acid (0.12/0.06-32/16); ampicillin (0.5-32, gram-negative panel, and 0.06-16, gram-positive panel); cefepime (0.5-32); ceftriaxone (0.06-64); ceftazidime (8-32); imipenem (0.06-16); linezolid (0.5-8); levofloxacin (0.008-8); minocycline (0.5-16); tigecycline (0.008-16); penicillin (0.06-8); piperacillin/tazobactam (0.06/4-128/4) and vancomycin (0.12-32). MIC interpretive criteria followed published guidelines established by the Clinical and Laboratory Standards Institute [8] and the recent US Food and Drug Administration package insert for tigecycline [9], where applicable.
- Escherichia coli*, *Klebsiella pneumoniae* and *Klebsiella oxytoca* were screened for ESBL activity when MIC results for ceftriaxone were >1 µg/ml using broth microdilution panels. ESBL activity was confirmed using the CLSI (2005) phenotypic confirmatory disk test (Oxoid, Ogdensburg, NY, USA) on Mueller-Hinton agar (Remel Inc., Lenexa, KS, USA) according to CLSI (2005) guidelines. ESBL presence was confirmed by testing the following antibiotic disks: cefotaxime (30-µg), cefotaxime/clavulanic acid (30/10-µg), ceftazidime (30-µg), and ceftazidime/clavulanic acid (30/10-µg). Antimicrobial disks were manufactured by Oxoid, Inc. (Ogdensburg, NY, USA). Mueller-Hinton agar used in testing was manufactured by Remel, Inc. (Lenexa, KS, USA). An organism was interpreted as containing an ESBL if there was an increase of >5 mm in the inhibition zone of the combination disk when compared to that of the cephalosporin alone.
- Quality controls (QC) were performed by each testing site on each day of testing using the corresponding ATCC control strains: *E. coli* ATCC 25922; *E. coli* ATCC 35218; *H. influenzae* ATCC 49766; *H. influenzae* ATCC 49247; *S. aureus* ATCC 29213; *Pseudomonas aeruginosa* ATCC 27853; *Enterococcus faecalis* ATCC 29212 and *S. pneumoniae* ATCC 49619. Results were included in the analysis only when corresponding QC isolates tested within the acceptable range according to CLSI (2005) guidelines [8].

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Results

The results are listed in the following table and figures.

Table 1. In vitro activity of tigecycline against selected pathogens by year of isolation.

Organism	n (04/05/06/07/08)	2004		2005		2006		2007		2008	
		MIC ₅₀	MIC ₉₀	MIC ₅₀	MIC ₉₀	MIC ₅₀	MIC ₉₀	MIC ₅₀	MIC ₉₀	MIC ₅₀	MIC ₉₀
<i>Acinetobacter</i> spp.	1496/1442/1983/2633/932	0.25	1	0.25	1	0.25	1	0.5	2	0.5	2
<i>A. baumannii</i>	1322/1321/1703/2333/834	0.5	1	0.5	1	0.5	2	0.5	2	0.5	2
<i>A. baumannii</i>	1123/845/1518/2822	0.06	0.25	0.12	0.5	0.12	0.5	0.12	0.5	0.12	1
<i>Enterobacteriaceae</i>	8579/9261/12720/16260/5156	0.5	1	0.5	1	0.5	1	0.5	1	0.5	1
<i>E. coli</i>	2551/2894/4063/5049/1617	0.12	0.25	0.12	0.25	0.12	0.5	0.12	0.5	0.25	0.5
<i>K. pneumoniae</i>	2086/2265/2981/3904/1266	0.5	2	0.5	2	0.5	2	0.5	2	0.5	2
<i>K. oxytoca</i>	506/493/678/938/269	0.25	1	0.25	1	0.25	1	0.25	1	0.25	1
<i>Klebsiella</i> spp.	2600/2766/3747/4911/1542	0.5	2	0.5	2	0.5	2	0.5	2	0.5	1
ESBL producers*	467/279/1007/1088/385	0.5	2	0.5	2	0.5	2	0.5	2	0.5	1
<i>E. aerogenes</i>	637/605/818/1037/390	0.5	1	0.5	1	0.5	2	0.5	1	0.5	2
<i>E. cloacae</i>	1704/1810/2480/3269/1056	0.5	2	0.5	2	0.5	2	0.5	2	0.5	2
<i>S. marcescens</i>	988/1071/1345/1722/538	1	2	1	2	1	2	1	2	1	2
<i>Enterococcus</i> spp.	1448/1732/2340/2959/963	0.06	0.12	0.06	0.12	0.06	0.12	0.12	0.25	0.12	0.25
<i>E. faecalis</i>	1096/1257/1580/2057/648	0.12	0.12	0.12	0.12	0.12	0.25	0.12	0.25	0.12	0.25
<i>E. faecium</i>	597/322/645/646/276	0.06	0.12	0.03	0.12	0.06	0.12	0.06	0.12	0.06	0.25
All VRE†	212/273/327/309/90	0.06	0.12	0.06	0.12	0.06	0.12	0.06	0.12	0.06	0.25
<i>S. aureus</i>	2541/2935/3599/4866/1520	0.12	0.25	0.12	0.25	0.12	0.25	0.12	0.25	0.12	0.25
MRSA‡	1174/1416/1541/2011/532	0.12	0.25	0.12	0.25	0.12	0.25	0.12	0.25	0.12	0.5
<i>S. pneumoniae</i>	1360/1504/1858/1945/279	0.03	0.12	0.03	0.06	0.03	0.06	0.03	0.12	0.06	0.12
(PISP)§	347/413/485/455/51	0.03	0.12	0.03	0.06	0.03	0.06	0.03	0.12	0.06	0.12
(PRSP)¶	166/201/245/309/30	0.03	0.12	0.03	0.06	0.03	0.06	0.03	0.12	0.06	0.12
<i>P. aeruginosa</i>	2077/2242/3105/4003/1279	8	>16	8	>16	8	>16	8	>16	8	>16

* ESBL = Extended spectrum beta-lactamase; includes *E. coli*, *K. pneumoniae* and *K. oxytoca*
 † VRE = Vancomycin resistant enterococci
 ‡ MRSA = Methicillin resistant *Staphylococcus aureus*
 § PISP = Penicillin intermediate *Streptococcus pneumoniae*; PRSP = Penicillin resistant *Streptococcus pneumoniae*

Figure 1. In vitro activity of tigecycline against 8,486 strains of *Acinetobacter* spp. by year of isolation.

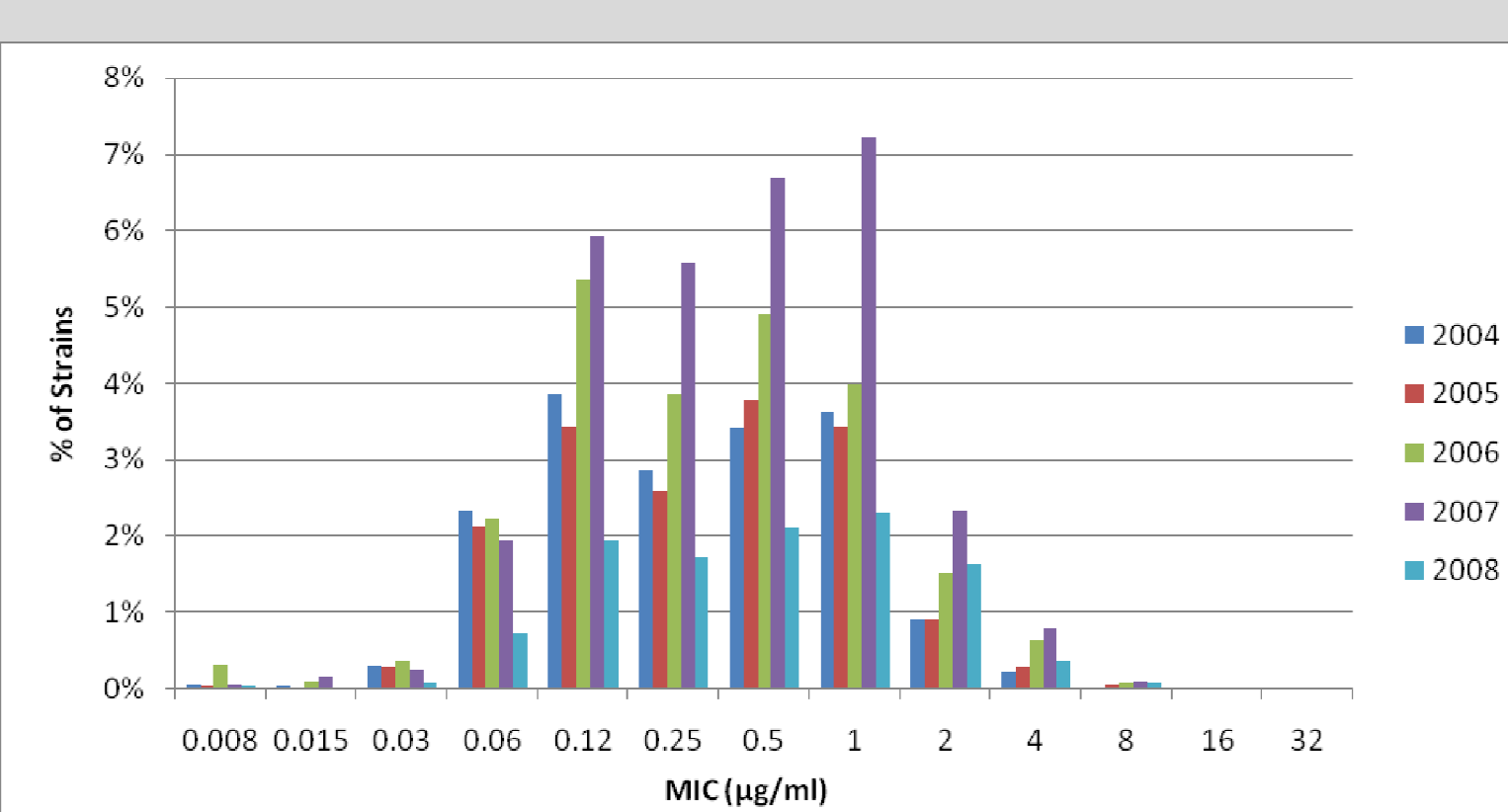


Figure 2. In vitro activity of tigecycline against 16,174 strains of *E. coli* by year of isolation.

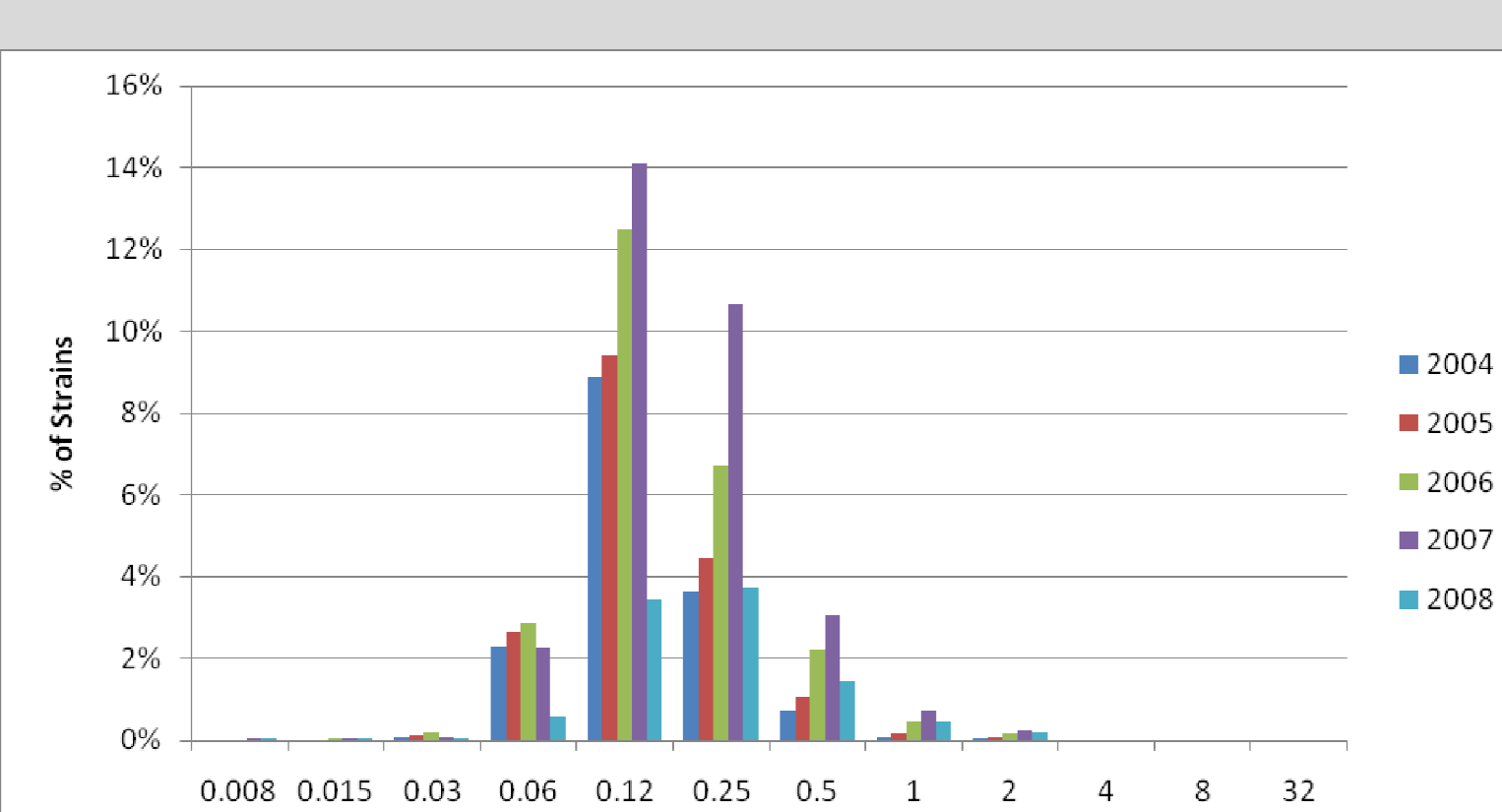


Figure 3. In vitro activity of tigecycline against 15,556 strains of *Klebsiella* spp. by year of isolation.

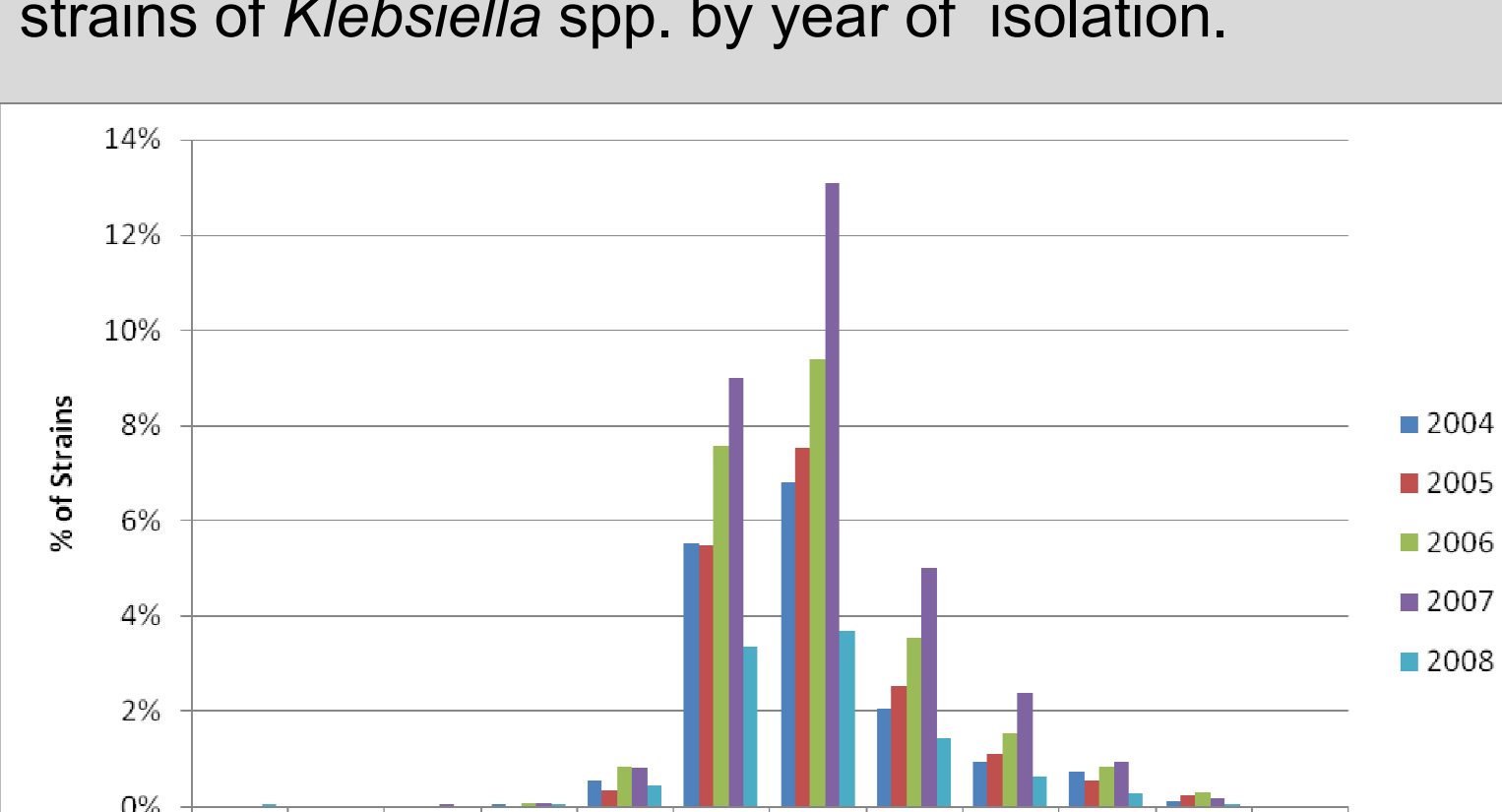


Figure 4. In vitro activity of tigecycline against 14,330 strains of *Enterobacter* spp. by year of isolation.

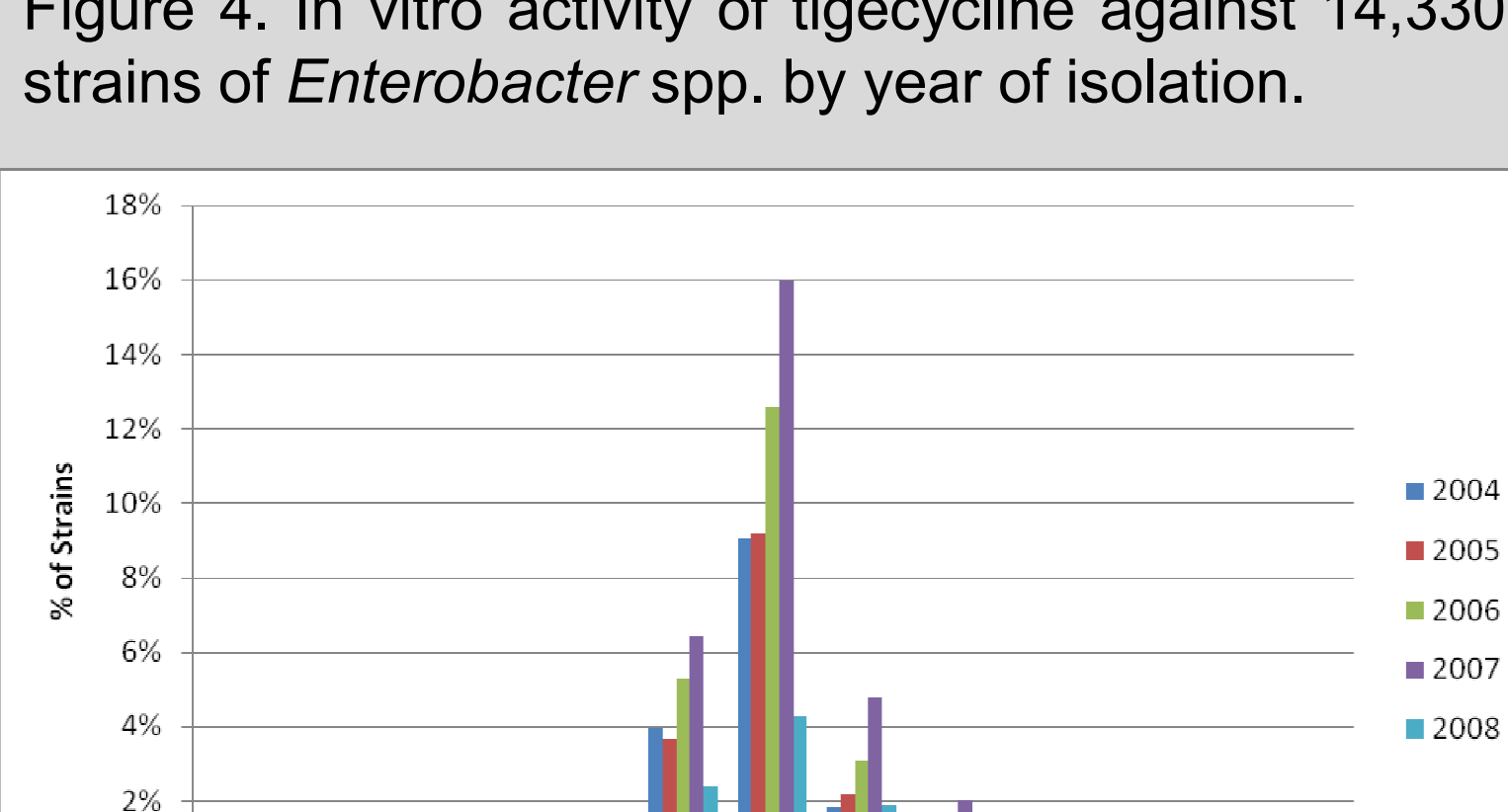


Figure 5. In vitro activity of tigecycline against 5,907 strains of *Serratia* spp. by year of isolation.

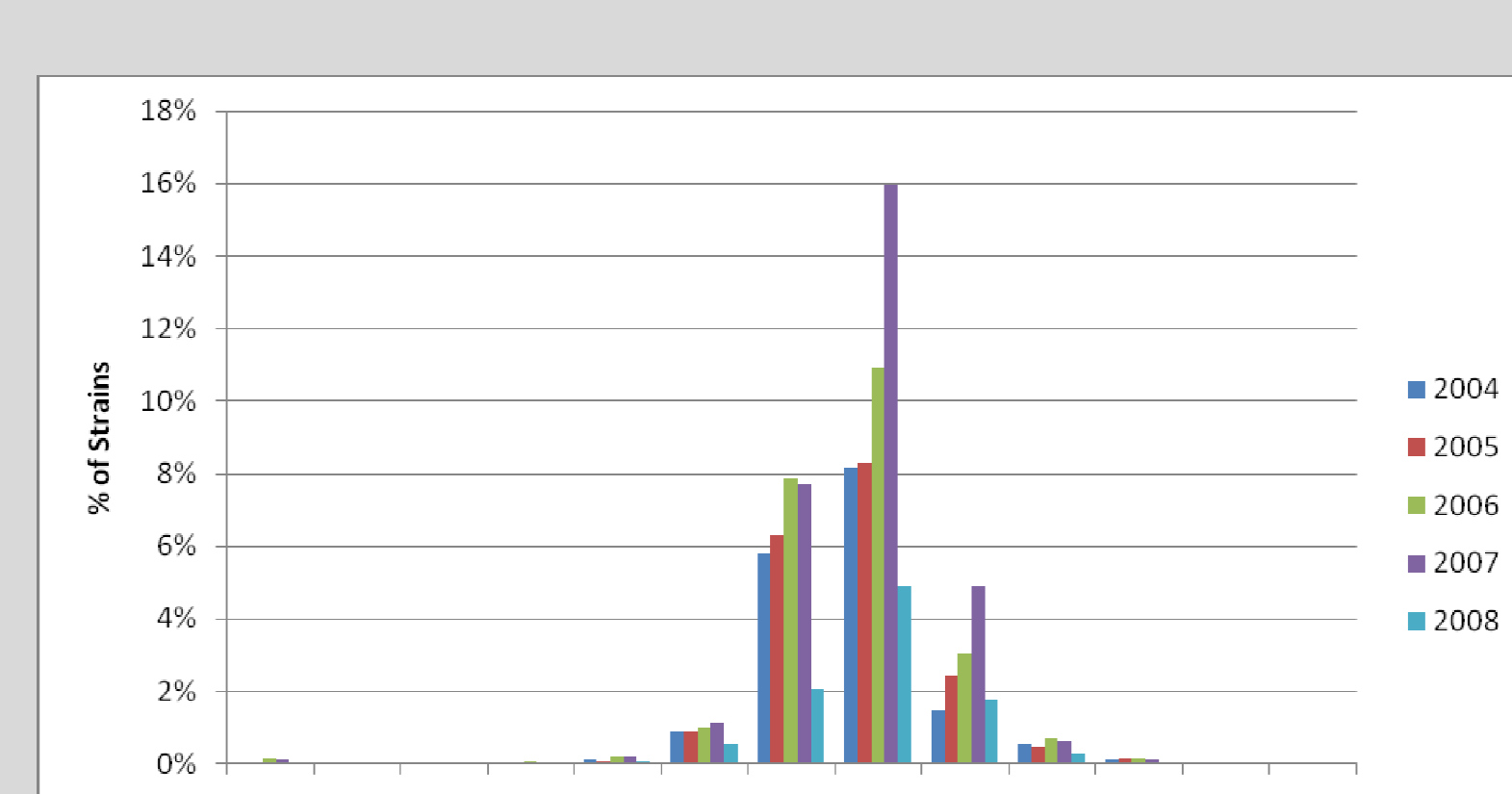


Figure 6. In vitro activity of tigecycline against 9,528 strains of *Enterococcus* spp. by year of isolation.

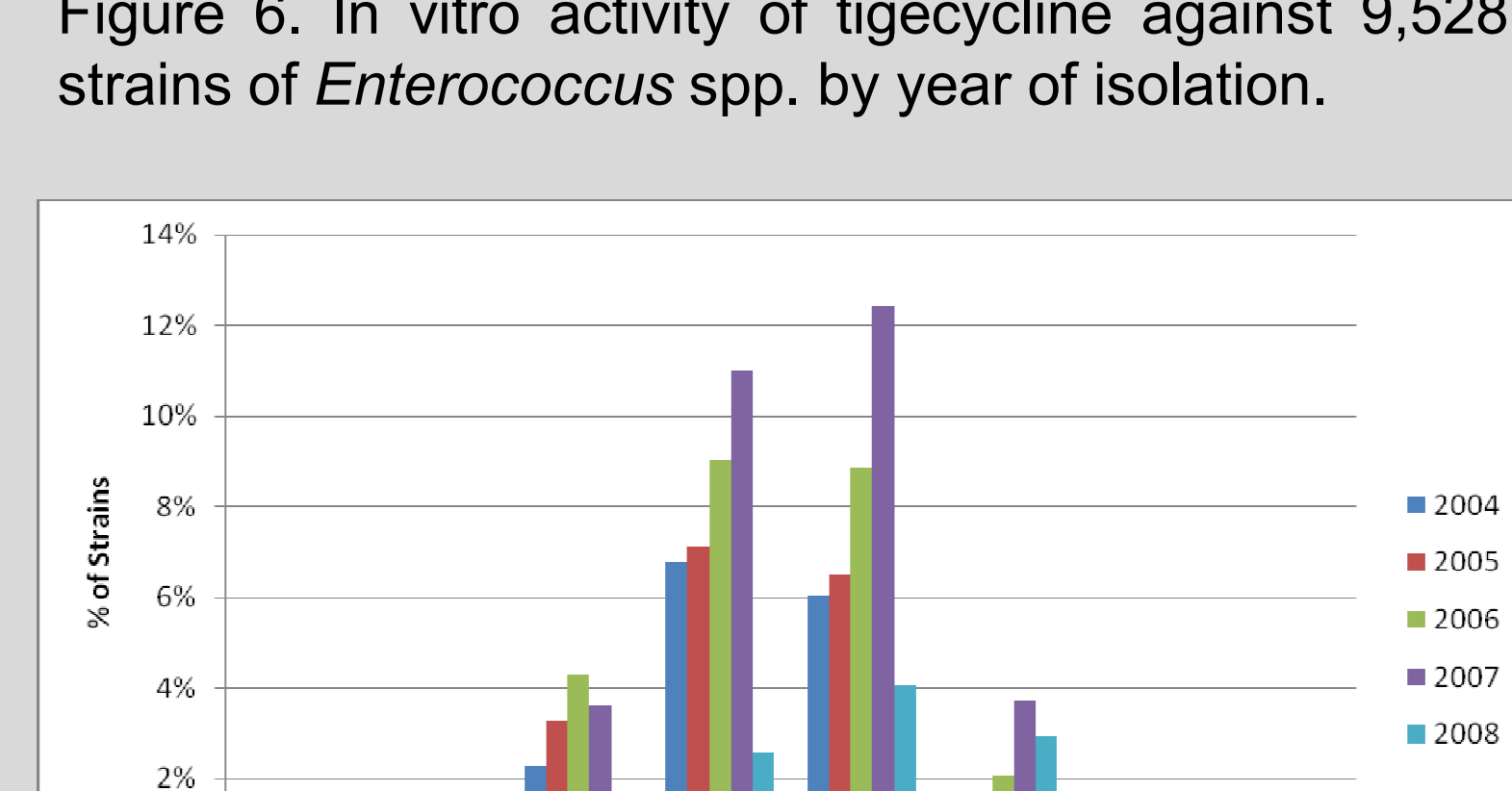


Figure 7. In vitro activity of tigecycline against 15,455 strains of *S. aureus* by year of isolation.

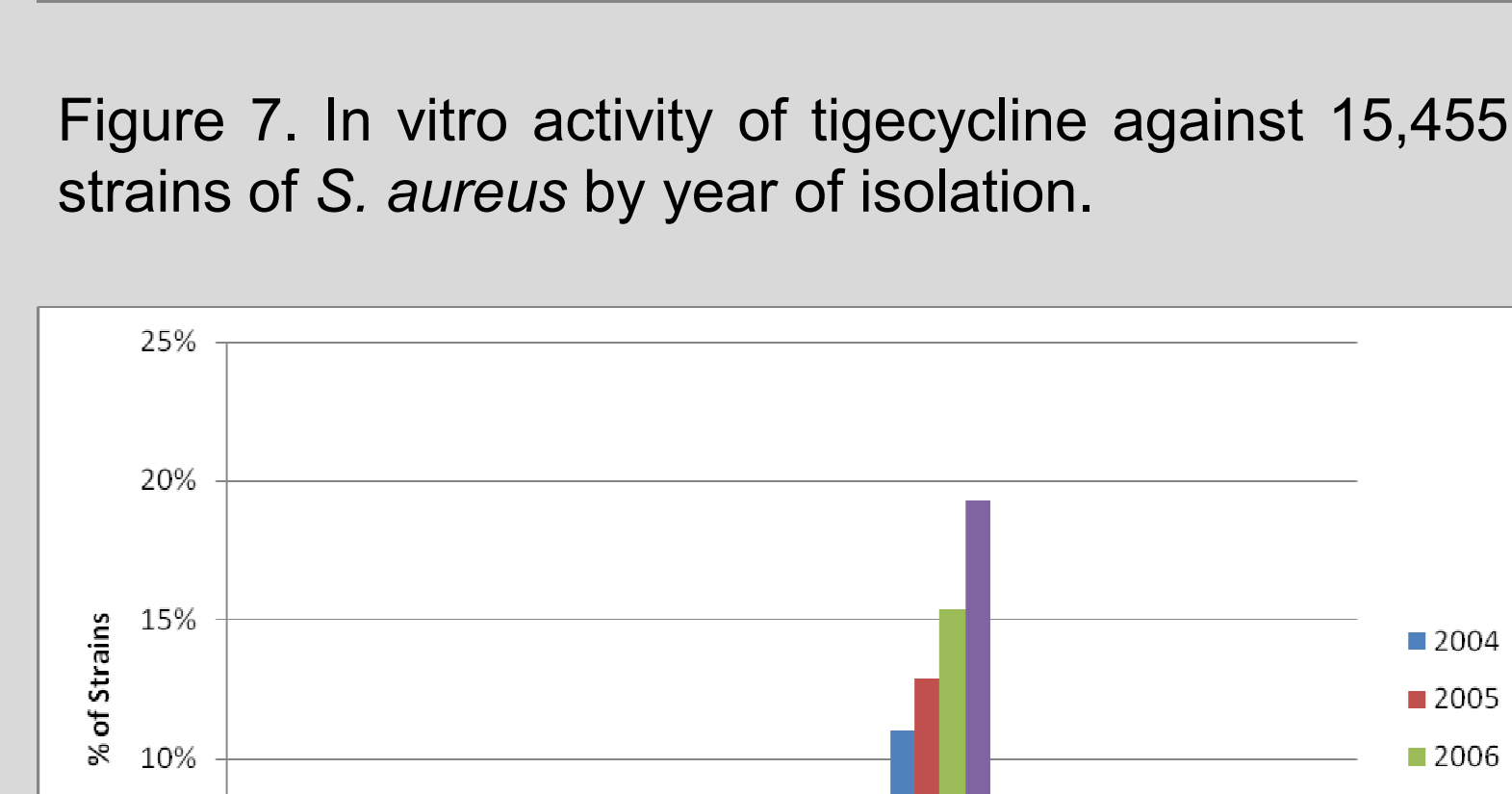
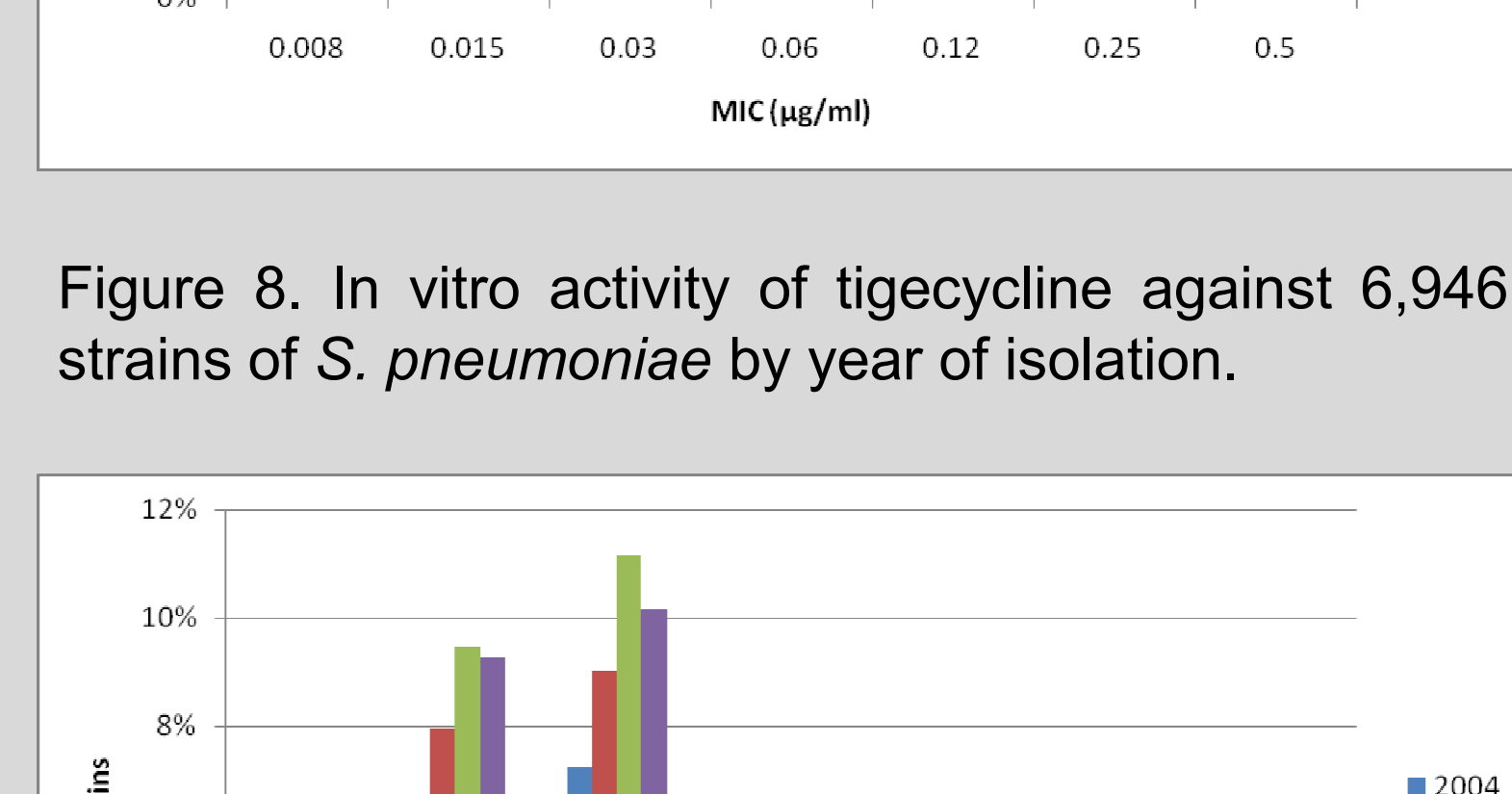


Figure 8. In vitro activity of tigecycline against 6,946 strains of *S. pneumoniae* by year of isolation.



Conclusions

- No MIC₅₀ or MIC₉₀ value for any organism group differed by more than a single dilution when comparing results from the years 2004 through 2008.
- During the five years covered by this analysis, tigecycline has fully retained its excellent activity against a broad spectrum of bacteria, including many strains resistant to various other antimicrobials.